FDA 510(k) Premarket Notification

K955651 MEDTRONIC SHERPA GUIDING CATHETER

CLEARED

Jan 22, 1996
41 days to decision

K955651 · Product code: **DQY** · Cardiovascular
Source: https://www.510kdatabase.net/k955651/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Dec 12, 1995
Decision date	Jan 22, 1996
Days to decision	41 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Medtronics Interventional Vascular**
Location	Danvers, MA, US
Contact	JOSEPH O MAGLIOZZI
510(k) history	21 submissions · 21 cleared · 1992-1999